KAEWOON FOR MEN- hyaluronic acid, allantoin gel
RNCARE

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

71237-003_V2

HYALURONIC ACID 2%

Allantoin 0.2%

Skin Protectant

Body Cleansing

Take an appropriate amount and massage onto body for about 60 seconds and rinse with water

In case of eye contact, immediately rinse with water

Stop use if unusual redness, swelling, soreness or irritation occur

Keep out of reach of children

Water, Polyquaternium-10 , Hydroxyethylcellulose , Black Strap Powder , Oleic Acid , Menthol , Lauryl Glucoside , Polysorbate 85 , 1,2-Hexanediol , Caprylic/Capric Triglyceride , Oryza Sativa (Rice) Bran Extract , Curcuma Longa (Turmeric Root) Extract, Vigna Radiata Seed Extract , Artemisia Princeps Extract , Arctium Lappa Root Extract , Hippophae Rhamnoides Extract , Hawthorn Crataegus Cuneata Extract Rheum Palmatum Root Extract , Bambusa Vulgaris Extract, Hamamelis Virginiana (Witch Hazel) Extract , Chamomilla Recutita (Matricaria) Extract , Panax Ginseng Root Extract , Propolis Extract , Sapindus Mukurossi Fruit Extract , Disodium EDTA , Diglycerin , Hydrolyzed Collagen , Butylene Glycol , Lauramidopropyl Betaine , Shea Butteramidopropyl Betaine , Capryl/Capramidopropyl Betaine , Sodium Cocoamphoacetate , Fragrance

KAEWOON FOR MEN

hyaluronic acid, allantoin gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71237-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYALURONIC ACID (UNII: S270N0TRQY) (HYALURONIC ACID - UNII:S270N0TRQY)	HYALURONIC ACID	20 mg in 1 mL
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	2 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
POLYQUATERNIUM-10 (1000 MPA.S AT 2%) (UNII: GMR4PEN8PK)	
HYDROXYETHYL CELLULOSE (100 MPA.S AT 2%) (UNII: R33S7TK2EP)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
POLYSORBATE 85 (UNII: A7F3N56197)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
RICE BRAN (UNII: R60QEP13IC)	
TURMERIC (UNII: 856YO1Z64F)	
MUNG BEAN (UNII: 1LIB31N73G)	
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)	
HIPPOPHAE RHAMNOIDES SEED (UNII: ZD5PJT4UZF)	
RHEUM PALMATUM ROOT (UNII: G025DAL7CE)	
BAMBUSA VULGARIS WHOLE (UNII: WCD45M1BSK)	
HAMAMELIS VIRGINIANA LEAF WATER (UNII: 8FP93ED6H2)	
MATRICARIA CHAMOMILLA ROOT (UNII: BTG5H50X7F)	
SAPINDUS MUKOROSSI FRUIT (UNII: 66H9NW427Y)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
LAURAMIDOPROPYL BETAINE (UNII: 23D6XVI233)	
SODIUM COCOAMPHOACETATE (UNII: W7Q5E87674)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71237-003-02	1 in 1 CARTON	11/21/2018	
1	NDC:71237-003-01	320 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/21/2018	

Labeler - RNCARE (694893883)

Registrant - RNCARE (694893883)

Establishment

Name	Address	ID/FEI	Business Operations
RNCARE		694893883	manufacture(71237-003)

Revised: 10/2021 RNCARE